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Datasheet for ABIN571086 anti-LPP antibody (Internal Region)

1 Image



Overview

Quantity:	100 µg
Target:	LPP
Binding Specificity:	Internal Region
Reactivity:	Human
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This LPP antibody is un-conjugated
Application:	Western Blotting (WB), ELISA

Product Details

Purpose:	LPP
Immunogen:	Peptide with sequence C-RNDSDPTYGQQGHP, from the internal region of the protein sequence according to NP_005569.1.
Sequence:	RNDSDPTYGQ QGHP
lsotype:	lgG
Cross-Reactivity:	Cow, Dog, Human
Purification:	Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.
Grade:	Verified

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Target Details		
Target:	LPP	
Alternative Name:	LPP (LPP Products)	
Background:	LPP, LIM domain containing preferred translocation partner in lipoma, LIM domain-containing	
	preferred translocation partner in lipoma, LIM protein, Lipoma-preferred partner	
Gene ID:	4026	
NCBI Accession:	NP_005569	
Application Details		
Application Notes:	Western Blot: Approx 85 kDa band observed in Human Placenta lysates and in lysates of cell	
	line HepG2 (calculated MW of 65.7 kDa according to NP_005569.1). This molecular weight is	
	routinely observed by other sources. Recommended concentration: 0.1-0.3 μ g/m	
	Peptide ELISA: antibody detection limit dilution 1:16000.	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	0.5 mg/mL	
Buffer:	Supplied at 0.5 mg/mL in Tris saline, 0.02 % sodium azide, pH 7.3 with 0.5 % bovine serum albumin.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which	
	should be handled by trained staff only.	
Handling Advice:	Minimize freezing and thawing.	
Storage:	-20 °C	

Storage Comment:Aliquot and store at -20°C, with minimal freeze/thawing. A working aliquot may be refrigeratedat 4°C for a few weeks and still remain viable.

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250kDa 150kDa	Western Blotting
100kDa 75kDa	Image 1. ABIN571086 (0.1µg/ml) staining of HepG2 lysate
50kDa	(35µg protein in RIPA buffer). Primary incubation was 1
37kDa	hour. Detected by chemiluminescence.
25kDa	
20kDa	
15kDa	

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